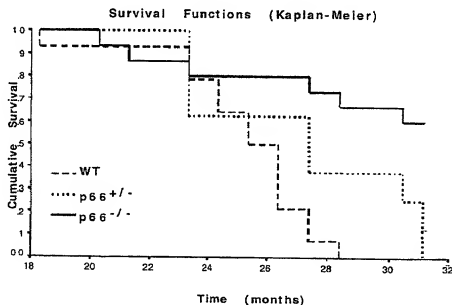




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(57) Abstract

It has been determined that i) p66^{shc} is serine phosphorylated upon UV treatment or oxidative damage; ii) the serine-phosphorylation of p66 by oxidative signals is mediated by Erk1 and p38, as shown both *in vivo* and *in vitro*; iii) ablation of p66^{shc} expression by homologous recombination enhances resistance to oxidative damage both *in vitro* and *in vivo*; iv) a serine-phosphorylation defective mutant of p66^{shc} is unable to restore a normal stress response in p66^{shc} targeted cells; v) mice carrying the p66^{shc} targeted mutation have prolonged lifespan.